AMENDMENT UNDER 37 C.F.R. § 1.111 Attorney Docket No.: Q101077

Application No.: 10/520,784

AMENDMENTS TO THE CLAIMS

This listing of claims will replace all prior versions and listings of claims in the application:

LISTING OF CLAIMS:

1. (Currently amended) A compound represented by the formula

$$\underbrace{ \left(\begin{array}{c} X \\ N \\ N \\ R^1 \end{array} \right) }_{R} Y \underbrace{ \left(\begin{array}{c} R^2 \\ N \\ R^3 \end{array} \right) }_{(I)}$$

wherein Ring A represents an optionally substituteda pyridine ring which may have one or two C₁₋₄ alkyl groups, X represents an electron attracting nitrile group, Y represents

-CH=CH- or -(CH₂)₂-(CH₃)₂, R¹ represents (1) a C₅₋₇ cycloalkyl group optionally fused with a benzene ring, (2) a C₇₋₁₉ aralkyl group, (3) a 5- or 6-membered heterocyclic ring-C₁₋₄ alkyl group or (4) a C₆₋₁₄ aryloxy-C₁₋₄ alkyl group, each of which may have 1 to 4 substituents selected from a halogen atom, a C₁₋₄ alkyl group, a mono-, di- or tri-halogeno-C₁₋₄ alkyl group and a C₁₋₄ alkyl group, and one of R² and R³ each independently represent is a hydrogen atom, an optionally substituted hydrocarbon group selected from the group consisting of a C₁₋₆-alkyl, C₃₋₁₀ eyeloalkyl, C₆₋₁₄ aryl and C₇₋₁₉ aralkyl or an optionally substituted heterocyclic group selected from the group consisting of tetrahydropyranyl, pyranyl and pyridyl or a C₁₋₄ alkyl group, and the other is a C₆₋₁₄ aryl group, a C₇₋₁₉ aralkyl group, a C₃₋₁₀ cycloalkyl group, a tetrahydropyranyl group, a pyranyl group or a pyridyl group or a C₁₋₆ alkyl group, a mono-, di- or tri- halogeno-C₁₋₄ substituents selected from a halogen atom, a C₁₋₄ alkyl group, a mono-, di- or tri- halogeno-C₁₋₄

AMENDMENT UNDER 37 C.F.R. § 1.111 Attorney Docket No.: Q101077

Application No.: 10/520,784

alkyl group, a C₁₋₄ alkoxy group, a C₁₋₄ alkoxy-carbonyl group, a cyano group, a C₁₋₄ alkyl-carbonylamino group and a hydroxyl group or a salt thereof.

- 2. (Canceled).
- (Canceled).
- (Canceled).
- 5. (Canceled).
- (Canceled).
- (Original) (2E)-3-(3-cyano-4,6-dimethyl-1-[(1S)-1,2,3,4-tetrahydronaphthalen-1-yll-1H-pyrrolo[2,3-b]pyridin-2-yl}-N-(3,4-dimethoxyphenyl)prop-2-enamide,
- (2E)-3-{3-cyano-4,6-dimethyl-1-[(1S)-1,2,3,4-tetrahydronaphthalen-1-yl]-1Hpyrrolo[2,3-b]pyridin-2-yl}-N-(3,4-dimethylphenyl)prop-2-enamide,
- (2E)-3-{3-cyano-4,6-dimethyl-1-[(1S)-1,2,3,4-tetrahydronaphthalen-1-yl]-1H-pyrrolo[2,3-b]pyridin-2-yl}-N-methyl-N-phenylprop-2-enamide,
- (2E)-3-{3-cyano-4,6-dimethyl-1-[(1S)-1,2,3,4-tetrahydronaphthalen-1-yl]-1Hpyrrolo[2,3-b]pyridin-2-yl}-N-(3-methylphenyl)prop-2-enamide,
- (2E)-3-{3-cyano-4,6-dimethyl-1-[(18)-1,2,3,4-tetrahydronaphthalen-1-yl]-1Hpyrrolo[2,3-b]pyridin-2-yl}-N-(4-hydroxy-3-methoxyphenyl)prop-2-enamide, or salts thereof.

AMENDMENT UNDER 37 C.F.R. § 1.111 Attorney Docket No.: Q101077

Application No.: 10/520,784

8. (Canceled).

9. (Currently amended) A medicine vanilloid receptor agonist comprising the

compound according to claim 1.

10. (Canceled).

11. (Currently amended) The vanilloid receptor agonist according to claim 10-9

which is for local administration.

12. (Currently amended) The vanilloid receptor agonist according to claim 10-9

which is an agent for treating overactive bladder.

13. (Currently amended) The vanilloid receptor agonist according to claim 10-9

which is an analgesic.

14. (Previously presented) A method of treating overactive bladder, comprising

administering to a mammal in need an effective amount of the compound according to claim 1.

15. (Previously presented) An analgesic method comprising administering to a

mammal in need an effective amount of the compound according to claim 1.

4

Attorney Docket No.: Q101077 AMENDMENT UNDER 37 C.F.R. § 1.111 Application No.: 10/520,784

(Canceled). 16.

(Canceled). 17.